ABSTRACT: Tuberculosis can involve essentially any ocular tissue. Tubercular Uveitis is difficult to diagnose because it may occur in patients without systemic manifestations of Tuberculosis. The diagnosis is therefore often presumptive, based on indirect evidence such as a positive skin test and negative findings for other causes of uveitis since a definitive diagnosis would require identification of the causative organism itself which is difficult to obtain. While the diagnosis of Central Serous Chorioretinopathy (CSCR) can be straightforward and accurate in majority of the cases, some unusual cases such as this require closer monitoring. A diagnosis of Tuberculosis should be in the backdrop of even typical appearing CSCR in India. A scar in the fundus of the other eye and a vague history of lymphadenopathy in childhood were the only suspicious factors in this case. Following investigations, a definitive diagnosis of choroidal tuberculosis was established, therapy instituted and resolution observed. The visual acuity recovered significantly after ATT and steroids.

KEY-WORDS: CSCR, OCT, Tubercular choroiditis.

INTRODUCTION: The two species causing TB in humans are the human strain M. tuberculosis, which is acquired by inhaling infected airborne droplets, and the bovine strain M. bovis, which is acquired by drinking unpasteurized milk from infected cattle. Ocular Tuberculosis can present with a myriad of symptoms and signs. Here we present a case report of Tubercular Choroiditis that presented atypically in a young, male patient.

CASE SUMMARY: A 32 year old male presented to eye department of PES medical college, Kuppam, rural Andhra Pradesh, in the month of June 2013 with complaints of sudden diminution of vision in left eye since that morning. He also gave history of seeing distorted images in the same eye. No history of trauma/redness/pain. He is a known hypertensive since two years, not on regular medication.

On questioning repeatedly history of hospitalization in childhood for bilateral inguinal swellings, the details of which were unavailable, was elicited.

Examination revealed normal anterior segments and pupillary reactions in both eyes. Vision was 6/6 in right eye (RE) and 6/60 improving to 6/18 with pinhole in left eye(LE). RE fundoscopy showed a well circumscribed circular scar of 250µ in the area inferonasal to the macula (Fig.1). In the LE, an elevated circular lesion with ring reflex was seen in the macula with absent foveal reflex. Initial provisional diagnosis of LE - CSCR was made.

On general physical examination, patient’s blood pressure was found to be 160/120 mm Hg. Patient was referred to physician for immediate systemic management of high BP.
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Patient was evaluated two days later, for ophthalmic investigations. Incidentally, deterioration of vision to counting fingers at 1m was noted along with increase in size of the lesion with hemorrhagic spots in LE (Fig.2)

Investigations:
Ophthalmic investigations: OCT revealed serous retinal detachment in left eye (Fig.3). FFA revealed hyper-fluorescence, increasing in size and intensity in late phase in the supero-temporal Macula (Fig.4). B-scan showed thickening of choroid (Fig.5), in the corresponding zone, with altered echoes.

Chest x-ray, Mantoux test, serology for Toxoplasma and HIV were done along with other routine laboratory investigations—Hb, TC, DC, ESR. Mantoux test read 28x30mm after 48hrs (Fig.6). Chest X-ray was normal (Fig.7). Other investigations were within normal range.

Treatment: A diagnosis of Tubercular Choroiditis was made and patient started on Category-1 Anti Tubercular Therapy (ATT), under the care of chest physician. Patient was reviewed every 3rd day. Systemic steroids were started 10 days after ATT.

Follow up: Improvement in vision in LE to 6/60 along with decrease in size of lesion with disappearance of hemorrhagic spots noted by 12th day of ATT (Fig.8). Two weeks later, with adjunctive oral corticosteroids, vision improved to 6/9 in the affected eye with fundus showing choroidal tubercle like picture (Fig. 9) After 4 weeks of initiating treatment, vision improved to 6/6.

DISCUSSION: Tuberculosis is known to cause a myriad of ophthalmic pathology. Approximately, 2 billion people are affected worldwide with tuberculosis, while only 10% of these people have clinical manifestations. The absence of pulmonary tuberculosis does not rule out ocular tuberculosis. Primary infection in the eye typically presents as corneal, conjunctival and scleral disease which is rare. Secondary ocular infection results from hematogenous spread of the organism or from a hypersensitivity reaction to extraocular infection.

Choroidal tubercles are the most common manifestation of intraocular tuberculosis. It may also present as multifocal progressive or diffuse choroiditis resembling serpiginous choroiditis.

Various criteria have been periodically updated for the diagnosis of the same. Though definitive diagnosis of ocular tuberculosis is established by the demonstration of the mycobacterium with the help of culture or DNA amplification from ocular samples, a presumptive diagnosis can be made based on ancillary testing after other causes of uveitis have been ruled out. A Mantoux test can be done if ocular tuberculosis is suspected. Greater than 5mm of induration after 48-72 hours is a positive result in a HIV patient. Greater than 10mm of induration is considered positive result in high-risk individuals, such as in people living in endemic areas, health-care workers and nursing home patients. Induration greater than 15mm is considered a positive result in all cases.

The present case had no signs of anterior segment or posterior segment inflammation which could have led to investigations for tuberculosis. A typical CSCR like picture with the backdrop of
systemic hypertension, was almost definitive. However, a small scar in the Right eye and vague history of lymph node biopsy in childhood led to further investigations to rule out tuberculosis.

CONCLUSION: Ocular Tuberculosis can have variable clinical manifestations. Patient's gender, age, his active life style and history of hypertension, initial clinical presentation and fundus findings led us to think in terms of CSCR. A high index of clinical suspicion is necessary especially in a country like ours where the disease is still rampant.

REFERENCES:

(Fig.1) Right Eye fundus picture of the patient

(Fig.2) LE fundus picture of the patient on second visit. The size of lesion has increased significantly with appearance of hemorrhagic spots.
(Fig.3) OCT of LE shows Retinal Detachment

(Fig.4) FFA shows blocked fluorescence inferotemporal to the disc in RE. LE shows late hyperfluorescence.

(Fig.5) B-Scan shows choroidal thickening.

(Fig.6) Mantoux test after 48 hours showing significant induration.
(Fig. 7) Chest X-ray PA view.

(Fig. 8) Fundus picture of LE after starting ATT. VA was 6/60

(Fig. 9) Fundus picture at subsequent follow up. VA improved to 6/9.

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